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⑤④ **Solvent for chromogenic dye-precursor material for pressure sensitive recording paper.**

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Description

The present invention relates to a solvent for a chromogenic dye-precursor material for a pressure-sensitive recording paper and to pressure-sensitive recording papers prepared by using the solvent.

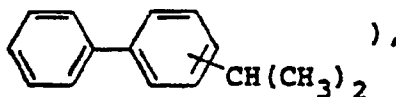
A pressure-sensitive recording paper sheet comprises a colour-development sheet prepared by coating microcapsules, in which a solution of a colourless electron-donating chromogenic dye-precursor material having a colouring reactivity has been encapsulated, onto a first support sheet and a colour-developer sheet prepared by coating a colour-developer which develops a colour on contact with the chromogenic dye-precursor material onto a second support sheet.

In recent years, the pressure-sensitive recording paper sheets have been widely used instead of carbon paper sheets and back-carbon copying paper sheets of the pigment type. It is necessary for the pressure-sensitive recording paper sheets that they exhibit excellent colour-development, stability for long term preservation and light resistance and that they are low in toxicity not to cause environmental pollution.

As the solvent which dissolves the chromogenic dye-precursor material in preparing the pressure-sensitive recording paper sheet, it is demanded that the solvent fulfills the following requirements.

- 1) To dissolve the chromogenic dye-precursor material to a high concentration.
- 2) Not to cause the decomposition and colour-development of the chromogenic dye-precursor material.
- 3) To show a considerably high boiling point, and not to evaporate in the thermal drying step and under high atmospheric temperature.
- 4) Not to reduce to water on encapsulating.
- 5) To show a high speed of colour-development and a high concentration of the developed colour as well as the high colour stability after colour-developing.
- 6) To be stable to light, heat and chemicals.
- 7) To show a low viscosity so that its flow out from the broken capsules is freely carried out.
- 8) To be substantially odorless.
- 9) To show a low toxicity to human body and to be safe.
- 10) To show a favorable biodegradability and not to cause environmental pollution.

As the solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet, which fulfills the above-mentioned requirements, several solvents have been proposed. For instance, (1) a mixed solvent comprising more than 65% by weight of isopropylbiphenyl (represented by the formula:



less than 25% by weight of polyisopropylbiphenyl and less than 10% by weight of biphenyl, which is used for preparing a solution of the chromogenic dye-precursor material to be contained in the capsules used for coating onto a sheet material for pressure-sensitive recording paper sheets (JP-B-54-37528(1979)), (2) hydrogenated terphenyl having at least 65% of aromaticity (hydrogenation rate of 35%), partially hydrogenated terphenyl, C₁₋₄-alkyl substituted terphenyl or a mixture thereof used as the solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet (JP-A-48-92112 (1973)), (3) a mixture of hydrogenated terphenyl (hydrogenation rate of not over 40%) and hexylbenzene or a mixture of C₁₀₋₁₆-alkylbenzene and C₇₋₁₀-alkylbenzene, which is a solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet (JP-A-No. 52-32922 (1977)) and (4) a solvent for the chromogenic dye-precursor material, comprising at least one of C₁₋₁₂ alkylated biphenyl or C₁₋₁₂-alkylated terphenyl, or a mixture of the alkylated-biphenyl or -terphenyl and other solvent [wherein the number of the alkyl group in the alkylated biphenyl is 1 to 4, that in the alkylated terphenyl is 1 to 6 and not less than two alkyls may be the same or different from each other] (GB-A-1352597).

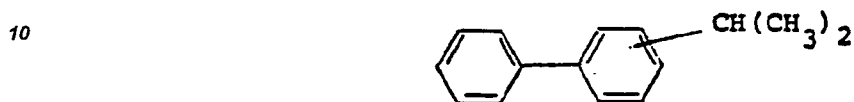
With the propagation of the pressure-sensitive recording paper sheets, the case wherein the pressure-sensitive recording paper sheets are used in cold districts of not more than -5°C in the ambient temperature or transported and preserved for a long time in the environment of about 40 to 50°C in the ambient temperature and of higher than about 80% in relative humidity has increased.

Particularly, in the outdoor facilities such as gasoline service stands, the pressure-sensitive recording paper sheets are used under the environment of lower than -5°C in winter.

Since in such an environment of low atmospheric temperature, 1) the solvent of the chromogenic dye-precursor material used in the pressure-sensitive recording paper sheet crystallizes, 2) it is necessary for a very long time in order to clearly develop or 3) the colour-developed image is very light in colour, if

developed, not to be deciphered, such a pressure-sensitive recording paper sheet is not to be put to practical use. Namely, it is demanded that an initial developing activity within 30 sec from the recording is at least 40%.

"Isopropylbiphenyl" disclosed in the JP-B-54-37528 (1979) as the solvent for the chromogenic dye-precursor material for the the pressure-sensitive recording paper sheet is a mixture of isomers represented by the formula :



15 wherein the isopropyl group occupies the o-, m- or p-position of the benzene ring of biphenyl.

The commercialized "isopropylbiphenyl" and the isopropylbiphenyl synthesized by Friedel-Crafts alkylation of biphenyl (refer to Industrial and Engineering Chemistry Product Research and Development, Vol. 8, 239 - 241, 1969) is a mixture of m-isomer and p-isomer containing a small amount of o-isomer. Such a mixed solvent exhales an offensive odor strongly and is not to be used as the solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet (refer to JP-B-50-14570 (1975)).

20 The problem of offensive odor occurs in the process for preparing the pressure-sensitive recording paper sheet when the pressure-sensitive recording paper sheet material prepared by coating the microcapsules containing the solution of the chromogenic dye-precursor onto the supporting sheet is cut after drying. Namely, when the pressure-sensitive recording paper sheet is cut at a relatively high temperature of 40 to 50°C, the solvent flowing out from the thus broken microcapsules gives a disagreeable impression to the operators. In addition, on the cases when the solvent adheres to clothes or hands in the cutting step of the pressure-sensitive recording paper sheet or in the handling of the solvent, the odor still remains even after washing the clothes or the hands with a cleanser to give a disagreeable impression. Furthermore, the pressure-sensitive recording paper sheets which have been subjected to recording are assembled and preserved in a storehouse for a relatively long time. In such occasion, the odor emitted from a large amount of the pressure-sensitive recording paper sheets which have been subjected to recording becomes to be the cause of disagreeable impression.

As has been shown above, the problem of the disagreeable odor concerning the pressure-sensitive recording paper sheets has been conspicuous in recent years.

35 As a result of the present inventors' studies for obtaining a solvent for the chromogenic dye-precursor material for the pressure-sensitive recording paper sheets, which is almost odorless, shows an excellent colour-development even at a low temperature, for instance, -5°C, and does not crystallize at such a low temperature of -5°C, it has been found that a solvent prepared by mixing p-monoisopropylbiphenyl with a hydrogenated terphenyl of a specified hydrogenation degree is almost odorless, does not crystallize at low temperatures of -5°C and fulfills all the above-mentioned requirements which are to be possessed by the solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet.

In a first aspect of the present invention, there is provided a substantially odorless solvent for the chromogenic dye-precursor material for a pressure-sensitive recording paper sheet, which solvent consists essentially of

- 45 (a) 50 to 80% by weight of p-monoisopropylbiphenyl or of a biphenyl mixture which is composed of not less than 80% by weight of p-monoisopropylbiphenyl, not more than 20% by weight of m-monoisopropylbiphenyl and not more than 10% by weight of diisopropylbiphenyl and which is substantially completely devoid of o-monoisopropylbiphenyl and
 (b) 50 to 20% by weight of hydrogenated terphenyl.

50 In a second aspect of the present invention, there is provided a process for the preparation of a substantially odorless solvent for a chromogenic dye-precursor material for a pressure-sensitive recording paper, which comprises mixing :

- 55 (a) 50 to 80% by weight of p-monoisopropylbiphenyl or of a biphenyl mixture which is composed of not less than 80% by weight of p-monoisopropylbiphenyl, not more than 20% by weight of m-monoisopropylbiphenyl and not more than 10% by weight of diisopropylbiphenyl and which is substantially completely devoid of o-monoisopropylbiphenyl, and
 (b) 50 to 20% by weight of hydrogenated terphenyl.

In a third aspect of the invention, there are provided microcapsules for a pressure-sensitive recording

paper, within which microcapsules is encapsulated a chromogenic dye-precursor material dissolved in a solvent according to the invention. Typically, the microcapsules comprise hydrophilic colloid walls containing a dye composition which is composed of the chromogenic dye-precursor material and the solvent of the invention. The microcapsules are prepared by a process comprising dissolving the chromogenic dye-precursor material in a solvent according to the invention and encapsulating said solution such as to form said microcapsules.

In a fourth aspect of the present invention, there is provided a pressure-sensitive recording paper wherein the microcapsules containing a solution of a chromogenic dye-precursor material are microcapsules according to the invention. The invention further provides a process for the preparation of a pressure-sensitive recording paper, which process comprises coating a first support sheet with microcapsules according to the invention to form a colour-development sheet, and providing said colour-development sheet with a colour-developer sheet comprising a second support sheet on which is coated a colour-developer such that the surface of said colour-development sheet provided with said microcapsules faces the surface of said colour-developer sheet provided with said colour developer.

The substantially odorless solvent for the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet according to the present invention (hereinafter referred to as the present solvent) consists essentially of 50 to 80% by weight of p-monoisopropylbiphenyl or a biphenyl mixture of not less than 80% by weight of p-monoisopropylbiphenyl, not more than 20% by weight of m-monoisopropylbiphenyl and not more than 10% by weight of diisopropylbiphenyl (hereinafter referred to as "p-monoisopropylbiphenyl" according to the present invention), the biphenyl mixture being substantially completely devoid of o-monoisopropylbiphenyl, and 50 to 20% by weight of hydrogenated terphenyl.

p-Monoisopropylbiphenyl contained in monoisopropylbiphenyl is almost odorless, excellent in dissolving the chromogenic dye-precursor material (determined at 20°C) but it melts at 11°C. In order to prevent the crystallization of p-monoisopropylbiphenyl at low temperatures such as -5°C, hydrogenated terphenyl is added thereto, and the thus obtained mixture is used as the solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet.

"p-Monoisopropylbiphenyl" according to the present invention may contain m-monoisopropylbiphenyl and diisopropylbiphenyl to the extent that they do not spoil the specificity of p-monoisopropylbiphenyl of almost odorless. Accordingly, "p-monoisopropylbiphenyl" according to the present invention is composed of not less than 80% by weight, preferably not less than 90% by weight of p-monoisopropylbiphenyl; not more than 20% by weight, preferably not more than 10% by weight, of m-mono-isopropylbiphenyl; and not more than 10% by weight, preferably not more than 5% by weight, of diisopropylbiphenyl, and does not contain o-monoisopropylbiphenyl.

"p-Monoisopropylbiphenyl" according to the present invention is produced, for instance, by the following processes.

(1) Biphenyl and propylene are reacted by heating to a temperature of 200 to 300°C, preferably 250 to 290°C, for 1 to 10 hours in the presence of silica-alumina catalyst. After the reaction is over, the catalyst is removed from the reaction mixture by filtration thereof, and the filtrate is subjected to rectification treatment, thereby obtaining "p-monoisopropylbiphenyl" according to the present invention, which contains not less than 80 % by weight of p-monoisopropylbiphenyl.

(2) Biphenyl and propylene are reacted by heating to a temperature of 200 to 300°C, preferably 220 to 290°C, for 1 to 10 hours in the presence of a zeolite catalyst. After the reaction is over, the catalyst is removed from the reaction mixture by filtration thereof, and the filtrate is subjected to rectification treatment, thereby obtaining "p-monoisopropylbiphenyl" according to the present invention, which contains not less than 80% by weight of p-monoisopropylbiphenyl.

(3) Biphenyl and propylene are reacted by heating to a temperature of 70 to 120°C, preferably 80 to 90°C, for 1 to 8 hours in the presence of aluminium chloride catalyst. After the reaction is over, the catalyst is removed from the reaction mixture, and the thus obtained organic layer is subjected to rectification treatment, thereby obtaining "p-monoisopropylbiphenyl" according to the present invention, which contains not less than 80% by weight of p-monoisopropylbiphenyl.

However, the production of "p-monoisopropylbiphenyl" according to the present invention is not limited to the abovementioned processes.

As has been described above, it is not necessary that "p-monoisopropylbiphenyl" according to the present invention is the single and pure compound of p-monoisopropylbiphenyl, however, any biphenyl mixture containing m-monoisopropylbiphenyl in an amount of over 20% by weight is not desirable because of the occurrence of problems of odor. In addition, it is necessary that o-monoisopropylbiphenyl is not contained in "p-monoisopropylbiphenyl" according to the present invention in view of odor and that the content of diisopropylbiphenyl is below 10% in view of the solubility of the chromogenic dye-precursor material in the

present solvent.

Another component of the present solvent, namely, "hydrogenated terphenyl" is composed of hydrogenated terphenyl having hydrogenation rate of not lower than 40%, and is produced by the following process.

5 Terphenyl is subjected to partial hydrogenation in the presence of a catalyst carrying at least one kind of metal selected from the group consisting of aluminum, palladium, platinum and rhodium at a temperature of 70 to 150°C and under a pressure of 10 to 100 kg/cm² to obtain the "partially" hydrogenated terphenyl to be used as "hydrogenated terphenyl" according to the present invention. However, "hydrogenated terphenyl" according to the present invention is not limited to the substance produced by the above-mentioned process.

10 "Hydrogenated terphenyl" according to the present invention is preferably the substance having hydrogenation rate of not lower than 40%, and in the case of below 40%, such a poorly hydrogenated terphenyl cannot attain the objective of the present invention.

As has been described, the present solvent is a mixture of 50 to 80% by weight of "p-monoisopropylbiphenyl" according to the present invention and 50 to 20% by weight of "hydrogenated terphenyl", and 15 in the case where the content of "p-monoisopropylbiphenyl" according to the present invention is over 80% by weight, crystals of p-monoisopropylbiphenyl precipitate from the solvent at low temperatures, for instance, -5°C and accordingly it is not desirable.

On the other hand, in the case where the content of "p-monoisopropylbiphenyl" according to the present invention is below 50%, since the initial colour-developing activity after 30 sec of recording at -5°C does 20 not attain the actually utilizable value and accordingly it is not desirable.

The heart of the pressure-sensitive recording paper sheet according to the present invention is characterized in that thy mixture of 50 to 80% by weight of "p-monoisopropylbiphenyl" according to the present invention and 50 to 20% by weight of hydrogenated terphenyl is used as a solvent of the chromogenic dye-precursor material. Accordingly, the present invention is not limited by the method of encapsulation, the 25 kinds of the chromogenic dye-precursor material, the colour-developer, the method of preparing the slurry of the above-mentioned materials and the method of coating the slurry onto the paper sheet material, namely, all the methods known by the persons skilled in the art can be applied in the present invention.

For instance, as the method of encapsulation, the method utilizing coacervation disclosed in US-A-2,800,457 and US-A-2,800,458 and the method by interfacial polymerization disclosed in GB-A-990,443 30 and US-A-3,287,154 are utilizable.

As the chromogenic dye-precursor material, compounds of triphenylmethanes, diphenylmethanes, xanthenes, thiazines and spiropyranes may be exemplified for the pressure-sensitive recording paper sheet.

Further, as the acidic substance used as the colour-developer, active clayish substances such as acidic clay, active clay, atapalgite, bentonite and zeolite or organoacidic substances such as phenol resin, acidic 35 reactive phenol-formaldehyde novolac resin and metal salts of aromatic organic acid may be exemplified.

The solvent for the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet according to the present invention is almost odorless and excellent in dissolving the chromogenic dye-precursor material, does not crystallize at lower temperature of -5°C and fulfills the necessary requirements which is to be provided by the above-mentioned solvent of the chromogenic dye-precursor for the pressure-sensitive recording paper sheet. 40

In addition, the initial colour-developing activity after 30 sec of recording at a low temperature of -5°C of the pressure-sensitive recording paper sheet according to the present invention is not lower than 40% and accordingly, the pressure-sensitive recording paper sheet according to the present invention can be applied to practical use even in cold districts.

45 The present invention will be concretely explained while referring to the non-limitative Examples, Comparative Examples and Reference Examples as follows.

EXAMPLE 1

50 Synthesis of p-monoisopropylbiphenyl

Into a 20-litre stainless steel autoclave provided with a heating apparatus, 12 kg of biphenyl and 1.5 kg of a silica-alumina catalyst (made by NIKKI Chemical Co., Ltd., X-632 HN) were introduced, and oxygen in the autoclave was substituted by nitrogen gas. Then the autoclave was heated to 70°C (inner temperature), and the stirring was commenced. From the same time, gaseous propylene was introduced into the autoclave from a propylene gas bomb to carry out the propylation of biphenyl in the autoclave. 55

Although the internal temperature of the autoclave raised slowly, the reaction temperature was maintained at about 280°C by controlling the heating apparatus. When the reduction of the weight of the pro-

pylene gas bomb became 3 kg, the supply of propylene was stopped and the reaction was continued further for one hour at the same temperature of 280°C, and then the autoclave was cooled.

After cooling the inner temperature to 40°C and taking the reaction mixture out from the autoclave, the catalyst was removed by filtration and the filtrate was subjected to rectification treatment while carrying out the analysis by gas-chromatography to obtain the object, p-monoisopropylbiphenyl. The thus obtained p-monoisopropylbiphenyl showed the following composition and physical properties.

Composition :

Biphenyl	0%
o-Monoisopropylbiphenyl	0%
m-Monoisopropylbiphenyl	6%
p-Monoisopropylbiphenyl	93%
Diisopropylbiphenyl	1%

Physical properties :

Specific gravity at 15°C	0.982
Refractive index at 25°C	1.5807
Boiling point	294°C
Temperature at which crystals precipitate	5°C

On subjecting the thus obtained p-monoisopropyl-biphenyl to a sensory test concerning the "yes" or "no" of the odor of 30 ml of the specimen thereof taken into a 100 ml wide mouth bottle by 20 men and 20 women of the panel, the number of person who answered "yes" was 2.

The above-mentioned result shows that the present solvent is excellent in odorlessness.

Preparation of the solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet

The present solvent was prepared by mixing 70% by weight of the thus produced p-monoisopropylbiphenyl and 30% by weight of odorless partially hydrogenated terphenyl (hydrogenation rate : 45%). Into 100 ml of the thus prepared solvent, 30 g of Crystal Violet Lactone (made by HODOGAYA Chemical Industry Co., Ltd.) (hereinafter referred to as CVL) were dissolved, and the concentration of CVL in the solution was determined in course of the time while keeping the solution in a thermostat at 20°C. The results are shown in Table 1.

Table 1

Time passed by (days)	1	7	14
Concentration of CVL (g/100 ml)	20	14	11

As are seen in Table 1, a state of high concentration of CVL was kept extremely stable for a long time. In addition, no crystal precipitated from the solution of CVL in the thus prepared solvent at -5°C.

EXAMPLE 2 :

Preparation of microcapsules

Microcapsules containing the mixed solvent prepared in Example 1 were prepared as follows.

A mixture of 630 g of melamine and 1620 g of an aqueous 37% solution of formaldehyde (hereinafter referred to as formalin) adjusted to pH of 9.0 by addition of an aqueous 2% solution of sodium hydroxide

was heated to 70°C, and after the dissolution of melamine into formalin, 2250 g of water were added to the mixture and the whole mixture was stirred for 3 min to obtain an aqueous solution of melamine-formaldehyde prepolymer.

Separately, a mixture of 600 g of urea and 1460 g of formalin adjusted to pH of 8.5 by triethanolamine was reacted at 70°C for 1 hour to obtain an aqueous solution of urea-formaldehyde prepolymer.

Separately, into a stirred mixture of 1620 g of formalin and 600 g of urea, triethanolamine was added to adjust the pH of the mixture to 8.8, and the mixture was reacted at 70°C for 30 min. Into 400 g of the thus obtained reaction mixture, 24 g of water and 30 g of tetraethylenepentamine were added and the pH of the thus prepared mixture was adjusted to 3 with an aqueous 15% solution of hydrochloric acid while stirring the mixture at 70°C. Since the pH of the mixture showed a reduction with the proceeding of the reaction, the pH of the mixture was readjusted to 3 by adding an aqueous 10% solution of sodium hydroxide, and then the reaction was continued at a reduced temperature of 55°C. When the viscosity of the reaction mixture became 200 cps, the reaction mixture was neutralized by adding the aqueous 10% solution of sodium hydroxide, and 4000 g of water were added to the thus neutralized reaction mixture to obtain an aqueous solution of water-soluble cationic urea resin.

After adjusting the pH of a mixture of 1000 g of the aqueous solution of melamine-formaldehyde prepolymer, 500 g of the aqueous solution of urea-formaldehyde prepolymer, 1580 g of the aqueous solution of water-soluble cationic urea resin, 620 g of water and 10 g of triethanolamine to 5.2 by the addition of an aqueous 10% solution of citric acid, 30 g of an aqueous 10% solution of a surfactant (made by KAO-Atlas Co., Ltd., NEOPELEX) was added to the mixture to obtain "A" liquid.

Separately, 500 g of Crystal Violet Lactone (a blue dye-precursor material made by HODOGAYA Chemical Industry Co., Ltd.) were dissolved in 9500 g of the mixed solvent prepared in Example 1 to obtain "B" liquid. 1000 ml of "B" liquid were homogenized into "A" liquid in a homogenizer so that the diameter of the thus formed particles of emulsion became from 2 to 8 µm. Thereafter, the thus formed emulsion was kept at 30°C while gently stirring and the pH thereof was adjusted to 3.6 by the addition of an aqueous 1% solution of citric acid. After stirring the thus adjusted emulsion for 1 hour, 2000 ml of water were added thereto.

After leaving the mixture further for 3 hours, an aqueous 20% solution of citric acid was added thereto to adjust the pH thereof to 3.0 and the mixture was stirred for 20 hours to obtain a slurry of microcapsules.

Preparation of the pressure-sensitive recording paper sheet

Into 600 ml of an aqueous 10% solution of polyvinyl alcohol (made by KURARE Co., Ltd., referred to as PVA), 300 g of the thus obtained microcapsules were added, and a dispersion of the microcapsules was prepared by stirring the mixture well.

The thus obtained aqueous dispersion was coated onto a paper sheet of 45 g/m² at a rate of 2.2 g of the microcapsules per m² of the paper sheet, and by superposing the thus treated paper sheet with a paper sheet on which a colour-developer comprising a condensate of p-phenylphenol and formaldehyde as the main colour-developer had been coating by a conventional method, a pressure-sensitive recording paper sheet was obtained.

After the colour-development of the thus obtained pressure-sensitive recording paper sheet by a typewriter made by Olivetti Co. in the ordinary environment and keeping the thus colour-developed paper sheet in a dark place for 24 hours, the concentration of the thus developed colour was measured by a reflex colour-densitometer made by MACBETH Co.

On the other hand, another pressure-sensitive recording paper prepared by the same process as above was subjected to colour-development in the environment of -5°C, and the concentration of the thus developed colour was measured by the same reflex colour densitometer from the time just after colour-development and the relative rate of colour-development was obtained in course of the time, in the case where the result of colour-development at ordinary temperature was appointed as 100, the results being shown in Table 2.

As will be seen in Table 2, the thus prepared pressure-sensitive recording paper sheet showed a sufficiently initial colour-developing activity of the thus obtained pressure-sensitive recording paper sheet after 30 sec of recording at a low temperature of -5°C.

Table 2

Time	30 sec	1 min	1 hour	24 hours
Rate of colour-development (%)	46	53	85	100

EXAMPLE 3 :

In the same manner as in Examples 1 and 2 except for using a solvent of the chromogenic dye-precursor material comprising 60 parts by weight of p-monoisopropylbiphenyl and 40 parts by weight of odorless partially hydrogenated terphenyl (hydrogenation rate : 45%), a solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet and a pressure-sensitive recording paper sheet were prepared, and the solubility of the chromogenic dye-precursor material and the initial colour-developing activity of the thus obtained pressure-sensitive recording paper sheet at low temperature of -5°C were measured. As the results, the solubility of the chromogenic dye-precursor material after 14 days at 20°C was 11.7 g/100 ml and the initial colour-developing activity at -5°C was 45%.

In addition, the thus obtained pressure-sensitive recording paper sheet was odorless, and no crystal precipitated in the thus obtained solvent at -5°C .

EXAMPLE 4 :

In the same manner as in Examples 1 and 2 except for using the solvent of the chromogenic dye-precursor material comprising 65 parts by weight of p-monoisopropylbiphenyl and 35 parts by weight of odorless partially hydrogenated terphenyl (hydrogenation rate : 50%), a solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet and a pressure-sensitive recording paper sheet were prepared, and the solubility of the chromogenic dye-precursor material and the initial colour-developing activity of the thus prepared pressure-sensitive recording paper sheet at a low temperature of -5°C were measured.

As the results, the solubility of the chromogenic dye-precursor material after 14 days at 20°C was 12.2 g/100 ml and the initial colour-developing activity at -5°C was 43%.

In addition, the thus obtained pressure-sensitive recording paper sheet was odorless, and no crystal precipitate in the solvent at -5°C .

COMPARATIVE EXAMPLE 1 :

Regarding p-monoisopropylbiphenyl used in Example 1, the solubility of the chromogenic dye-precursor material for pressure-sensitive recording paper sheet was examined. As a result, the solubility of the chromogenic dye-precursor material after 14 days was 9.0 g/100 ml at 20°C . Namely, although p-monoisopropylbiphenyl showed an excellent solubility of the chromogenic dye-precursor material in the same extent as the solvent prepared in Example 1, precipitation of crystals was observed at low temperatures of around 0°C .

COMPARATIVE EXAMPLE 2 :

In the same manner as in Example 1, a solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet was prepared except for mixing 90 parts by weight of p-monoisopropylbiphenyl and 10 parts by weight of odorless partially hydrogenated terphenyl (hydrogenation rate : 45%). The solubility of the thus prepared solvent of the chromogenic dye-precursor material after 14 days was 9.6 g/100 ml at 20°C . Namely, although the solvent showed an excellent solubility of the chromogenic dye-precursor material in the same extent as the solvent according to the present invention, precipitation

of crystals was observed at low temperatures around 0°C.

COMPARATIVE EXAMPLE 3 :

5 In the same manner as in Examples 1 and 2 except for using the solvent of the chromogenic dye-precursor material comprising 40 parts by weight of p-monoisopropylbiphenyl and 60 parts by weight of odorless partially hydrogenated terphenyl (hydrogenation rate : 45%), a solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet and a pressure-sensitive recording paper sheet were prepared, and the initial colour-developing activity of the thus prepared pressure-sensitive recording paper sheet at -5°C was examined. The initial colour-developing activity at -5°C was 33%.

COMPARATIVE EXAMPLE 4 :

15 In the same manner as in Example 2 except for using only the odorless partially hydrogenated terphenyl (hydrogenation rate : 45%) as the solvent of the chromogenic dye precursor material, a pressure-sensitive recording paper sheet was prepared, and the initial colour-developing activity thereof at low temperatures was examined. The initial colour-developing activity at -5°C was less than 10%.

COMPARATIVE EXAMPLE 5 :

20 In order to prevent the precipitation of crystals from p-monoisopropylbiphenyl at low temperatures, 1-xylyl-1-phenylethane was admixed with p-monoisopropylbiphenyl as follow.

Namely, 30 parts by weight of 1-xylyl-1-phenylethane and 70 parts by weight of p-monoisopropylbiphenyl were mixed together to prepare a solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet, and the thus prepared solvent was subjected to a sensory test concerning odor by 40 persons of the panel. As a result, all 40 persons answered that the solvent had an odor, and in addition, 23 persons complained an unpleasant odoriferousness.

Accordingly, the solvent prepared by admixing 1-xylyl-1-phenylethane with p-monoisopropylbiphenyl is not suitable as the solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheets.

EXAMPLE 5 :

Synthesis of p-monoisopropylbiphenyl mixture

35 Into a 2-litre stainless steel autoclave provided with a heating apparatus, 0.8 kg of biphenyl and 0.1 kg of zeolite Y-type catalyst (made by TOYO SODA Co., Ltd., TSZ-330 HUA, Dry : 300°C for 3 hours) were introduced, and oxygen in the autoclave was substituted by nitrogen gas. Then the autoclave was heated to 70°C (inner temperature), and the stirring was commenced. From the same time, propylene was introduced into the autoclave from a propylene gas-bomb to carry out the propylation of biphenyl in the autoclave.

40 Although the internal temperature of the autoclave raised slowly, the reaction temperature was maintained at about 270°C by controlling the heating apparatus. When the reduction of the weight of the propylene gas-bomb became 0.2 kg, the supply of propylene was stopped and the reaction was continued further for one hour at the same temperature of 270°C, and then the autoclave was cooled.

45 After cooling the inner temperature to 40°C and taking the reaction mixture out from the autoclave, the catalyst was removed by filtration and the filtrate was subjected to rectification treatment while carrying out the analysis by gaschromatography to obtain the object, p-monoisopropylbiphenyl mixture. The thus obtained p-monoisopropylbiphenyl mixture showed the following composition and physical properties.

50 Composition :

biphenyl	0%
o-monoisopropylbiphenyl	0%
m-monoisopropylbiphenyl	2%
p-monoisopropylbiphenyl	91%
55 diisopropylbiphenyl	7%

Physical properties :

Specific gravity at 15°C 0.988

Refractive index at 25°C	1.582
Boiling point	292°C
Temperature at which crystals precipitate	4°C

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On subjecting the thus obtained p-monoisopropylbiphenyl mixture to a sensory-test concerning the "yes" or "no" of the odor of 30 ml of the specimen thereof taken into a 100 ml wide mouth bottle by 20 men and 20 women of the panel, the number of person who answered "yes" was 3.

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The above-mentioned result shows that the thus obtained p-monoisopropylbiphenyl mixture is excellent in odorlessness.

Preparation of the pressure-sensitive recording paper sheet

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In the same manner as in Examples 1 and 2 except for using the thus obtained p-monoisopropylbiphenyl mixture, a solvent of the chromogenic dye-precursor material for pressure-sensitive recording paper sheet and a pressure-sensitive recording paper sheet were prepared, and the solubility of the chromogenic dye-precursor material and initial colour developing activity of the thus obtained pressure-sensitive recording paper sheet at low temperature were examined. As the result, the solubility of the chromogenic dye-precursor material after 14 days at 20°C was 13.8 g/100 ml and the initial colour-development at -5°C was 45%.

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Further, the thus prepared pressure-sensitive recording paper sheet was odorless and no precipitation of crystals was observed at -5°C.

EXAMPLE 6 :

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Synthesis of p-monoisopropylbiphenyl mixture

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Into a 1-litre glass autoclave provided with a heating apparatus, 0.5 kg of biphenyl and 0.014 kg of aluminium chloride catalyst were introduced, and oxygen in the autoclave was substituted by nitrogen gas. Then the autoclave was heated to 70°C (inner temperature), and the stirring was commenced. From the same time, propylene was introduced into the autoclave from a propylene gas-bomb to carry out the propylation of biphenyl in the autoclave.

35

Although the internal temperature of the autoclave raised slowly, the reaction temperature was maintained at about 90°C by controlling the heating apparatus. Propylene was supplied into the autoclave for 6 hours and when the reduction of the weight of the propylene gas-bomb became 0.165 kg, the supply of propylene was stopped, and then the autoclave was cooled.

40

After cooling the inner temperature to 40°C and taking the reaction mixture out from the autoclave, the catalyst was removed and the thus obtained organic layer was subjected to rectification treatment while carrying out the analysis by gas-chromatography to obtain the object, p-monoisopropylbiphenyl mixture. The thus obtained p-monoisopropylbiphenyl mixture showed the following composition and physical properties.

Composition :

biphenyl	0%
o-monoisopropylbiphenyl	0%
45 m-monoisopropylbiphenyl	4%
p-monoisopropylbiphenyl	95%
diisopropylbiphenyl	1%

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Physical properties :

Specific gravity at 15°C	0.986
Refractive index at 25°C	1.585
Boiling point	292°C
Temperature at which crystals precipitate	4°C
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On subjecting the thus obtained p-monoisopropylbiphenyl mixture to a sensory-test concerning the "yes" or "no" of the odor of 30 ml of the specimen thereof taken into a 100 ml wide mouth bottle by 20 men

and 20 women of the panel, the number of person who answered "yes" was 8.

The above-mentioned result shows that the thus obtained p-monoisopropylbiphenyl mixture is excellent in odorlessness.

5 Preparation of the pressure-sensitive recording paper sheet

In the same manner as in Examples 1 and 2 except for using the thus obtained p-monoisopropylbiphenyl mixture, a solvent of the chromogenic dye-precursor material for the pressure-sensitive recording paper sheet and a pressure-sensitive recording paper sheet were prepared, and the solubility of the chromogenic dye-precursor material and initial colour-developing activity of the thus obtained pressure-sensitive recording paper sheet at low temperature were examined. As the result, the solubility of the chromogenic dye-precursor material after 14 days at 20°C was 13.5 g/100 ml and the initial colour-development at -5°C was 44%. Further, the thus prepared pressure-sensitive recording paper sheet was odorless and no precipitation of crystals was observed at -5°C.

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REFERENCE EXAMPLE 1 :

Synthesis of m-monoisopropylbiphenyl

20 Into a 20-litre stainless-steel autoclave provided with a heating apparatus, 12 kg of biphenyl and 1.5 kg of a silica-alumina catalyst (made by NIKKI Chemical Co., Ltd., X-632 HN) were introduced, and after substituting oxygen in the autoclave by nitrogen gas, the content of the autoclave was heated.

When the inner temperature of the autoclave raised to 70°C, the stirring was commenced and at the same time, gaseous propylene was introduced into the autoclave to carry out the propylation.

25 Although the inner temperature of the autoclave raised slowly, the reaction temperature was maintained at about 260°C by controlling the heating apparatus.

When the reduction of the weight of the bomb became 3 kg, the supply of propylene was stopped, and after continuing the reaction for one hour at the same temperature of 260°C, the autoclave was cooled.

30 After cooling the autoclave to 40°C, the liquid reaction mixture was taken out from the autoclave and the catalyst was removed from the reaction mixture by filtration. The filtrate was subjected to rectification treatment while analyzing the distillate by gas-chromatography to obtain m-monoisopropylbiphenyl of a purity of 93%.

35 As a result of subjecting 100 ml of the thus obtained m-monoisopropylbiphenyl taken in a 300 ml-wide mouth bottle to a sensory-test concerning odors, 36 persons of all 40 persons of the panel answered "yes", and 13 persons of 36 persons complained an unpleasant odoriferousness.

REFERENCE EXAMPLE 2 :

Synthesis of o-monoisopropylbiphenyl

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Into a 20-litre stainless-steel autoclave provided with a heating apparatus, 12 kg of biphenyl and 1.5 kg of a silica-alumina catalyst (made by NIKKI Chemical Co., Ltd., X-632 HN) were introduced, and after substituting oxygen in the autoclave by nitrogen gas, the content of the autoclave was heated.

45 When the inner temperature of the autoclave raised to 70°C, the stirring was commenced and at the same time, gaseous propylene was introduced into the autoclave to carry out the propylation.

Although the inner temperature of the autoclave raised slowly, the reaction temperature was maintained at about 190°C by controlling the heating apparatus.

When the reduction of the weight of the bomb became 3 kg, the supply of propylene was stopped, and after continuing the reaction for one hour at the same temperature of 190°C, the autoclave was cooled.

50 After cooling the autoclave to 40°C, the liquid reaction mixture was taken out from the autoclave and the catalyst was removed from the reaction mixture by filtration. The filtrate was subjected to rectification treatment while analyzing the distillate by gas-chromatography to obtain o-monoisopropylbiphenyl of a purity of 83%.

55 As a result of subjecting 100 ml of the thus obtained m-monoisopropylbiphenyl taken in a 300 ml-wide mouth bottle to a sensory test concerning odors, all 40 persons of the panel answered "yes", and 26 persons complained an unpleasant odoriferousness.

Claims**Claims for the following Contracting States : CH, DE and LI**

- 5 1. A substantially odorless solvent for a chromogenic dye-precursor material for a pressure-sensitive recording paper, which solvent consists essentially of :
- (a) 50 to 80% by weight of p-monoisopropylbiphenyl or of a biphenyl mixture which is composed of not less than 80% by weight of p-monoisopropylbiphenyl, not more than 20% by weight of m-monoisopropylbiphenyl and not more than 10% by weight of diisopropylbiphenyl and which is substantially completely devoid of o-monoisopropylbiphenyl, and
- 10 (b) 50 to 20% by weight of hydrogenated terphenyl.
2. A solvent according to claim 1, wherein said biphenyl mixture consists essentially of not less than 90% by weight of p-monoisopropylbiphenyl, not more than 10% by weight of m-monoisopropylbiphenyl and not more than 5% by weight of diisopropylbiphenyl.
- 15 3. A process for the preparation of a substantially odorless solvent for a chromogenic dye-precursor material for a pressure-sensitive recording paper, which comprises mixing :
- (a) 50 to 80% by weight of p-monoisopropylbiphenyl or of a biphenyl mixture which is composed of not less than 80% by weight of p-monoisopropylbiphenyl, not more than 20% by weight of m-monoisopropylbiphenyl and not more than 10% by weight of diisopropylbiphenyl and which is substantially completely devoid of o-monoisopropylbiphenyl, and
- 20 (b) 50 to 20% by weight of hydrogenated terphenyl.
4. A process according to claim 3, wherein said p-monoisopropylbiphenyl or biphenyl mixture has been produced by reacting biphenyl with propylene at a temperature of 200 to 300°C for 1 to 10 hours in the presence of a silica-alumina catalyst or a zeolite catalyst and subjecting the obtained reaction mixture to rectification treatment.
- 25 5. A process according to claim 3, wherein said p-monoisopropylbiphenyl or biphenyl mixture has been produced by reacting biphenyl with propylene at a temperature of 70 to 120°C for 1 to 8 hours in the presence of an aluminium chloride catalyst and subjecting the obtained reaction mixture to rectification treatment.
6. Microcapsules for a pressure-sensitive recording paper, within which microcapsules is encapsulated a chromogenic dye-precursor material dissolved in a solvent as claimed in claim 1 or 2.
- 30 7. A process for the preparation of microcapsules for a pressure-sensitive recording paper, which process comprises dissolving a chromogenic dye-precursor material in a solvent as claimed in claim 1 or 2 and which has been provided by a process as claimed in any one of claims 3 to 5, and encapsulating said solution such as to form said microcapsules.
- 35 8. A pressure-sensitive recording paper wherein the microcapsules containing a solution of a chromogenic dye-precursor material are microcapsules as claimed in claim 6.
9. A process for the preparation of a pressure-sensitive recording paper, which process comprises coating a first support sheet with microcapsules as claimed in claim 6 and which have been prepared by a process as claimed in claim 7 to form a colour-development sheet, and providing said colour-development sheet with a colour-developer sheet comprising a second support sheet on which is coated a colour-developer such that the surface of said colour-development sheet provided with said microcapsules faces the surface of said colour-developer sheet provided with said colour-developer.
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Claim for the following Contracting State : ES

- 45 1. A process for the preparation of a pressure-sensitive recording paper, said process being characterised by comprising :
- (a) mixing (1) 50 to 80% by weight of p-monoisopropylbiphenyl or of a biphenyl mixture composed of not less than 80% by weight of p-monoisopropylbiphenyl, not more than 20% by weight of m-monoisopropylbiphenyl and not more than 10% by weight of diisopropylbiphenyl and which is substantially completely devoid of o-monoisopropylbiphenyl, and (2) 50 to 20% by weight of hydrogenated terphenyl, thereby obtaining a substantially odorless solvent for a chromogenic dye-precursor material ;
- 50 (b) dissolving a chromogenic dye-precursor material in the solvent obtained in step (a) and encapsulating said solution such as to form microcapsules ; and
- 55 (c) coating a first support sheet with the microcapsules obtained in step (b), to form a colour-development sheet and providing said colour-development sheet with a colour-developer sheet comprising a second support sheet on which is coated a colour-developer such that the surface of said colour-development sheet provided with said microcapsules faces the surface of said colour-developer sheet

provided with said colour-developer.

2. A process according to claim 1, wherein said biphenyl mixture consists of not less than 90% by weight of p-monoisopropylbiphenyl, not more than 10% by weight of m-monoisopropylbiphenyl and not more than 5% by weight of diisopropylbiphenyl.

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Ansprüche

Patentansprüche für folgende Vertragsstaaten : CH, DE und LI

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1. Im wesentlichen geruchloses Lösungsmittel für ein chromogenes Farbvorgläufermaterial für ein druckempfindliches aufzeichnungspapier, wobei das Lösungsmittel im wesentlichen besteht aus :

(a) 50 bis 80 Gew.% p-Monoisopropylbiphenyl oder einer Biphenylmischung, welche aus nicht weniger als 80 Gew.% p-Monoisopropylbiphenyl, nicht mehr als 20 Gew.% m-Monoisopropylbiphenyl und nicht mehr als 10 Gew.% Diisopropylbiphenyl besteht, und welche im wesentlichen vollständig frei von o-Monoisopropylbiphenyl ist ; und

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(b) 50 bis 20 Gew.% hydriertes Terphenyl.

2. Lösungsmittel gemäss Anspruch 1, wobei die Biphenylmischung im wesentlichen aus nicht weniger als 90 Gew.% p-Monoisopropylbiphenyl, nicht mehr als 10 Gew.% m-Monoisopropylbiphenyl und nicht mehr als 5 Gew.% Diisopropylbiphenyl besteht.

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3. Verfahren zum Herstellen eines im wesentlichen geruchlosen Lösungsmittels für ein chromogenes Farbvorgläufermaterial für ein druckempfindliches Aufzeichnungspapier, welches das Mischen von :

(a) 50 bis 80 Gew.% p-Monoisopropylbiphenyl oder einer Biphenylmischung, welche aus nicht weniger als 80 Gew.% p-Monoisopropylbiphenyl, nicht mehr als 20 Gew.% m-Monoisopropylbiphenyl und nicht mehr als 10 Gew.% Diisopropylbiphenyl zusammengesetzt ist und welche im wesentlichen vollständig frei ist von o-Monoisopropylbiphenyl ; und

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(b) 50 bis 20 Gew.% hydriertes Terphenyl umfasst.

4. Verfahren gemäss Anspruch 3, wobei das p-Monoisopropylbiphenyl oder die Biphenylmischung durch Umsetzen von Biphenyl mit Propylen bei einer Temperatur von 200 bis 300°C für 1 bis 10 Stunden in Anwesenheit eines Siliziumoxid-Aluminiumoxid-Katalysators oder eines Zeolit-Katalysators und durch Unterwerfen der so erhaltenen Reaktionsmischung unter eine Rektifizierbehandlung hergestellt worden war.

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5. Verfahren gemäss Anspruch 3, wobei das p-Monoisopropylbiphenyl oder die Biphenylmischung durch Umsetzen von Biphenyl mit Propylen bei einer Temperatur von 70 bis 120°C für 1 bis 8 Stunden in Anwesenheit eines Aluminiumchlorid-Katalysators und durch Unterwerfen der erhaltenen Reaktionsmischung unter eine Rektifizierbehandlung hergestellt worden war.

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6. Mikrokapseln für ein druckempfindliches Aufzeichnungsmaterial, wobei in den Mikrokapseln ein in einem Lösungsmittel gemäss Ansprüchen 1 oder 2 aufgelöstes chromogenes Farbvorgläufermaterial eingekapselt ist.

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7. Verfahren zum Herstellen von Mikrokapseln für ein druckempfindliches Aufzeichnungsmaterial, wobei das Verfahren das Auflösen eines chromogenen Farbvorgläufermaterials in einem Lösungsmittel gemäss Ansprüchen 1 oder 2 und welches durch ein Verfahren gemäss Ansprüchen 3 bis 5 zur Verfügung gestellt worden war, und das Einkapseln der Lösung unter Erhalt der Mikrokapseln umfasst.

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8. Druckempfindliches Aufzeichnungspapier, worin die eine Lösung eines chromogenen Farbvorgläufermaterials enthaltenden Mikrokapseln gemäss Anspruch 6 sind.

9. Verfahren zum Herstellen eines druckempfindlichen Aufzeichnungspapiers, welches das Beschichten eines ersten Trägerblattes mit Mikrokapseln gemäss Anspruch 6 und welche durch ein Verfahren gemäss Anspruch 7 hergestellt worden waren, unter Bildung eines Farbentwicklungsblattes, und das Versehen dieses Farbentwicklungsblattes mit einem Farbentwicklerblatt, welches ein zweites Trägerblatt umfasst, auf welches ein Farbentwickler schichtförmig aufgetragen ist, so dass die Oberfläche des Farbentwicklungsblattes, das mit den Mikrokapseln versehen ist, die Oberfläche des Farbentwicklerblattes bedeckt, das mit dem Farbentwickler versehen ist, umfasst.

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Patentanspruch für folgenden Vertragsstaat : ES

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1. Verfahren zum Herstellen eines druckempfindlichen Aufzeichnungsblattes, wobei das Verfahren dadurch gekennzeichnet, dass es umfasst :

(a) Mischen (1) von 50 bis 80 Gew.% p-Monoisopropylbiphenyl oder einer Biphenylmischung, die aus

nicht weniger als 80 Gew.% p-Monoisopropylbiphenyl, nicht mehr als 20 Gew.% m-Monoisopropylbiphenyl und nicht mehr als 10 Gew.% Diisopropylbiphenyl zusammengesetzt ist, und welche im wesentlichen vollständig frei ist von o-Monoisopropylbiphenyl, und (2) 50 bis 20 Gew.% hydriertes Terphenyl, wobei man ein im wesentlichen geruchsfreies Lösungsmittel für ein chromogenes Farbvorläufermaterial erhält ;

(b) Auflösen eines chromogenen Farbvorläufermaterials in dem nach Schritt (a) erhaltenen Lösungsmittel und das Einkapseln der Lösung, so dass Mikrokapseln entstehen ; und

(c) Beschichten eines ersten Trägerblattes mit in Schritt (b) erhaltenen Mikrokapseln unter Ausbildung eines Farbentwicklungsblattes und Versehen dieses Farbentwicklungsblattes mit einem Farbentwicklerblatt, welches ein zweites Trägerblatt umfasst, auf welches ein Farbentwickler in der Weise aufgebracht ist, dass die Oberfläche des Farbentwicklungsblattes, die mit den Mikrokapseln versehen ist, die Oberfläche des Farbentwicklerblattes, das mit dem Farbentwickler versehen ist, bedeckt.

2. Verfahren gemäss Anspruch 1, worin die Biphenylmischung aus nicht weniger als 90 Gew.% p-Monoisopropylbiphenyl, nicht mehr als 10 Gew.% m-Monoisopropylbiphenyl und nicht mehr als 5 Gew.% Diisopropylbiphenyl besteht.

Revendications

Revendications pour les états contractants : CH, DE et LI

1. Solvant essentiellement inodore de matériau précurseur de colorant chromogène pour papier d'enregistrement sensible à la pression, ledit solvant consistant essentiellement de :

(a) 50 à 80% en poids de p-monoisopropylbiphényle ou d'un mélange de biphényles qui est constitué de pas moins de 80% en poids de p-monoisopropylbiphényle, pas plus de 20% en poids de m-monoisopropylbiphényle et pas plus de 10% en poids de diisopropylbiphényle et qui est essentiellement dépourvu de o-monoisopropylbiphényle, et

(b) 50 à 20% en poids de terphényle hydrogéné.

2. Solvant suivant la revendication 1, dans lequel ledit mélange de biphényles consiste essentiellement de pas moins de 90% en poids de p-monoisopropylbiphényle, pas plus de 10% en poids de m-monoisopropylbiphényle et pas plus de 5% en poids de diisopropylbiphényle.

3. Procédé pour la préparation d'un solvant essentiellement inodore de matériau précurseur de colorant chromogène pour papier d'enregistrement sensible à la pression, ledit procédé comprenant le mélange de :

(a) 50 à 80% en poids de p-monoisopropylbiphényle ou d'un mélange de biphényles qui est constitué de pas moins de 80% en poids de p-monoisopropylbiphényle, pas plus de 20% en poids de m-monoisopropylbiphényle et pas plus de 10% en poids de diisopropylbiphényle et qui est essentiellement dépourvu de o-monoisopropylbiphényle, et

(b) 50 à 20% en poids de terphényle hydrogéné.

4. Procédé suivant la revendication 3, dans lequel ledit p-monoisopropylbiphényle ou ledit mélange de biphényles a été préparé par réaction du biphényle avec le propylène à une température de 200 à 300°C pendant 1 à 10 heures en présence d'un catalyseur silice-alumine ou zéolite puis soumission du mélange réactionnel ainsi obtenu à un traitement de rectification.

5. Procédé suivant la revendication 3, dans lequel ledit p-monoisopropylbiphényle ou ledit mélange de biphényles a été préparé par réaction du biphényle avec le propylène à une température de 70 à 120°C pendant 1 à 8 heures en présence de chlorure d'aluminium catalyseur puis soumission du mélange réactionnel ainsi obtenu à un traitement de rectification.

6. Microcapsules pour papier d'enregistrement sensible à la pression, dans lesquelles est encapsulé un matériau précurseur de colorant chromogène dissout dans un solvant tel que revendiqué dans la revendication 1 ou 2.

7. Procédé pour la préparation de microcapsules pour papier d'enregistrement sensible à la pression, ledit procédé comprenant la dissolution d'un matériau précurseur de colorant chromogène dans un solvant tel que revendiqué dans la revendication 1 ou 2 et qui a été obtenu selon le procédé de l'une quelconque des revendications 3-5, et l'encapsulation de ladite solution de façon à former lesdites microcapsules.

8. Papier d'enregistrement sensible à la pression dans lequel les microcapsules contenant une solution d'un matériau précurseur de colorant chromogène sont des microcapsules telles que revendiquées dans la revendication 6.

9. Procédé pour la préparation d'un papier d'enregistrement sensible à la pression, ledit procédé comprenant le revêtement d'une première feuille support avec des microcapsules telles que revendiquées

dans la revendication 6 et qui ont été préparées selon un procédé tel que revendiqué dans la revendication 7 pour former une feuille de développement chromogène, et l'application de ladite feuille de développement chromogène à une feuille de révélation chromogène comprenant une seconde feuille support sur laquelle est revêtu un révélateur chromogène, de telle façon que la surface de ladite feuille de développement chromogène pourvue desdites microcapsules soit située en regard de la surface de ladite feuille de révélation chromogène pourvue dudit révélateur chromogène.

Revendication pour l'état contractant : ES

1. Procédé pour la préparation d'un papier d'enregistrement sensible à la pression, ledit procédé comprenant :

(a) le mélange de : (1) 50 à 80% en poids de p-monoisopropylbiphényle ou d'un mélange de biphényles qui est constitué de pas moins de 80% en poids de p-monoisopropylbiphényle pas plus de 20% en poids de m-monoisopropylbiphényle et pas plus de 10% en poids de diisopropylbiphényle et qui est essentiellement dépourvu de o-monoisopropylbiphényle, et (2) 50 à 20% en poids de terphényle hydrogéné, de façon à obtenir un solvant essentiellement inodore de matériau précurseur de colorant chromogène; (b) la dissolution d'un matériau précurseur de colorant chromogène dans le solvant obtenu au stade (a) et l'encapsulation de la solution résultante de façon à former des microcapsules ; et

(c) le revêtement d'une première feuille support avec des microcapsules obtenues au stade (b), pour former une feuille de développement chromogène, et l'application de ladite feuille de développement chromogène à une feuille de révélation chromogène comprenant une seconde feuille support sur laquelle est revêtu un révélateur chromogène, de telle façon que la surface de ladite feuille de développement chromogène pourvue desdites microcapsules soit située en regard de la surface de ladite feuille de révélation chromogène pourvue dudit révélateur chromogène.

2. Procédé suivant la revendication 1, dans lequel ledit mélange de biphényles consiste essentiellement de pas moins de 90% en poids de p-monoisopropylbiphényle, pas plus de 10% en poids de m-monoisopropylbiphényle et pas plus de 5% en poids de diisopropylbiphényle.